

SHAMPOO COMPOSITIONS AND METHOD

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Abstract of WO9805296

An aqueous conditioning shampoo composition comprising, in addition to water: i) at least one surfactant chosen from anionic, nonionic, zwitterionic or amphoteric surfactants or mixtures thereof; ii) emulsified particles of an insoluble, nonvolatile silicone; iii) a soluble cationic hair conditioning polymer having a cationic charge density of about +3.0 meq/gram or less, in which the emulsified particles of insoluble, nonvolatile silicone are incorporated into the shampoo composition as a preformed aqueous emulsion having an average silicone particle size in the emulsion and in the shampoo composition of from 2 to 30 microns.

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- 1 -

SHAMPOO COMPOSITIONS AND METHOD

5 Field of the Invention

This invention relates to shampoo compositions, and more particularly to shampoo compositions containing emulsified particles of silicone, which compositions condition the hair
10 leaving it softer and more manageable.

Background and Prior Art

The use of silicones as conditioning agents in cosmetic
15 formulations is well known and widely documented in the patent literature. Generally, dispersed droplets of the silicone oil are suspended in the composition, which is then applied to the hair to deposit the silicone material on the hair shaft.

20

A typical method of silicone shampoo manufacture is disclosed in WO 92/10162. Essentially, the silicone material is emulsified directly into the shampoo by an *in situ* hot process, in which the complete shampoo mixture
25 incorporating the silicone is mixed thoroughly at elevated temperature, pumped through a high shear mill and then cooled. The silicone can be dispersed in a first process stage with anionic surfactant and fatty alcohol to form a premix. The premix is then mixed with the remaining
30 materials of the shampoo, pumped through a high shear mill, and cooled to obtain the final composition.

- 2 -

A disadvantage associated with an *in situ* hot process such as is described in WO 92/10162 is that factory handling of viscous silicone oil is difficult in the context of a full shampoo manufacturing operation.

5

A further disadvantage is that special equipment is normally needed to control silicone particle size during manufacture.

GB 2 170 216 A discloses a similar process, in which the full shampoo composition incorporating insoluble, non-

10 volatile silicone is sheared with a high shear mixer until the silicone particles are on average less than 2 microns in diameter. The particle size distribution is then said to be from about 2 to about 55 microns.

15 In order to solve the above mentioned problems with *in situ* hot processing of silicone, the alternative of incorporating the silicone as a preformed aqueous emulsion has been proposed. Such a method has the consequences that the silicone is incorporated with a predeterminable, 20 controllable particle size distribution. The silicone is insoluble and remains emulsified in the fully formulated shampoo composition, and thus the step of high shear processing of the silicone within the fully formulated shampoo composition is not required. This also makes 25 manufacture of the compositions easier.

A typical method for incorporating insoluble, non-volatile silicone materials into a conditioning shampoo is disclosed in US 5,085,087 in which such materials are incorporated in

30 the shampoo composition as a pre-formed aqueous emulsion of average particle size less than 2 microns. All the ingredients are mixed in a simple hot or cold process in which the average particle size of the silicone material in

- 3 -

the emulsion remains the same in the final shampoo composition. Preferably, this size is from 0.01 to 1 micron, e.g. 0.4 micron.

5 EP 0 529 883 A1 discloses hair shampoo compositions made by an equivalent method and comprising microemulsified particles of silicone having a particle size of 0.15 microns or less, e.g., 0.036 microns. Reducing the silicone particle size still further in this way is said to improve stability,
10 optical properties and conditioning performance.

The shampoos of US 5,085,087 and EP 0 529 883 A1 require the presence of a cationic polymer to deposit the silicone efficiently from the formulation onto the hair. A problem
15 encountered with these small particle size silicone/cationic polymer formulations is that they can give an undesirable sensory feel, typically manifest as a perception of "slippiness" and/or heaviness on dry hair.

20 We have now found that the utilisation of insoluble, non-volatile silicone, in the form of an aqueous, preformed emulsion of large particle size in a surfactant-based shampoo composition will impart conditioning benefit to the hair without the dry hair sensory negatives associated with
25 the prior art compositions discussed above.

Surprisingly and advantageously, we have found that in compositions of our invention, the presence of cationic polymer selectively enhances the wet properties of the
30 shampoo, e.g., wet feel and ease of wet comb - whilst reducing the ease of dry combing. In contrast, we found that the presence of cationic polymer in prior art formulations incorporating cationic polymer and smaller

- 4 -

particle size pre-formed silicone emulsions did not give a selective increase in ease of wet combing. Ease of dry combing was increased to at least the same extent.

- 5 The selective enhancement of wet properties such as ease of wet combing observed with compositions of the present invention is of benefit to the consumer. This is because consumers need to detangle their hair easily when wet, but when it is dry, if the hair slips through the comb too easily then it can be difficult to put into style.
- 10

Summary of the Invention

- 15 The invention provides an aqueous shampoo composition comprising, in addition to water:
- i) at least one surfactant chosen from anionic, nonionic, zwitterionic or amphoteric surfactants or mixtures thereof;
- 20
- ii) emulsified particles of an insoluble, nonvolatile silicone;
- iii) a cationic hair conditioning polymer having a cationic charge density of about +3.0 meq/gram or less,
- 25
- in which the emulsified particles of insoluble, non-volatile silicone are incorporated into the shampoo composition as a preformed aqueous emulsion having an average silicone particle size in the emulsion and in the shampoo composition of from 2 to 30 microns.
- 30

- 5 -

In another aspect, the invention provides a method of making such a shampoo composition, by mixing together water, the surfactant, the cationic conditioning polymer and a preformed aqueous emulsion of the silicone, wherein the 5 silicone in the emulsion and in the shampoo composition has an average particle size of from 2 to 30 microns.

Detailed Description of the Invention

10 i) Surfactant

The composition according to the invention comprises a surfactant chosen from anionic, nonionic, zwitterionic or amphoteric surfactants or mixtures thereof.

15

Suitable anionic surfactants include the alkyl sulphates, alkyl ether sulphates, alkaryl sulphonates, alkanoyl isethionates, alkyl succinates, alkyl sulphosuccinates, N-alkoyl sarcosinates, alkyl phosphates, alkyl ether phosphates, alkyl ether carboxylates, and alpha-olefin sulphonates, especially their sodium, magnesium ammonium and mono-, di- and triethanolamine salts. The alkyl and acyl groups generally contain from 8 to 18 carbon atoms and may be unsaturated. The alkyl ether sulphates, alkyl ether phosphates and alkyl ether carboxylates may contain from one to 10 ethylene oxide or propylene oxide units per molecule, and preferably contain 2 to 3 ethylene oxide units per molecule.

30 Examples of suitable anionic surfactants include sodium oleyl succinate, ammonium lauryl sulphosuccinate, ammonium lauryl sulphate, sodium dodecylbenzene sulphonate, triethanolamine dodecylbenzene sulphonate, sodium cocoyl

- 6 -

isethionate, sodium lauroyl isethionate and sodium N-lauryl sarcosinate. The most preferred anionic surfactants are sodium lauryl sulphate, triethanolamine lauryl sulphate, triethanolamine monolauryl phosphate, sodium lauryl ether 5 sulphate 1EO, 2EO and 3EO, ammonium lauryl sulphate and ammonium lauryl ether sulphate 1EO, 2EO and 3EO.

Nonionic surfactants suitable for use in compositions of the invention may include condensation products of aliphatic (C₈-10 C₁₈) primary or secondary linear or branched chain alcohols or phenols with alkylene oxides, usually ethylene oxide and generally having from 6 to 30 ethylene oxide groups. Other suitable nonionics include mono- or di-alkyl alkanolamides. Example include coco mono- or di- ethanolamide and coco 15 mono-isopropanolamide.

Amphoteric and zwitterionic surfactants suitable for use in compositions of the invention may include alkyl amine oxides, alkyl betaines, alkyl amidopropyl betaines, alkyl 20 sulphobetaines (sultaines), alkyl glycimates, alkyl carboxyglycimates, alkyl amphopropionates, alkylamphoglycimates alkyl amidopropyl hydroxysultaines, acyl taurates and acyl glutamates, wherein the alkyl and acyl groups have from 8 to 19 carbon atoms. Examples 25 include lauryl amine oxide, cocodimethyl sulphopropyl betaine and preferably lauryl betaine, cocamidopropyl betaine and sodium cocamphopropionate.

The surfactants are present in shampoo compositions of the 30 invention in an amount of from 0.1 to 50% by weight, preferably from 0.5 to 30% by weight.

- 7 -

Generally, the surfactants are present in shampoo compositions of the invention in an amount of from 0.1 to 50%, preferably from 5 to 30%, more preferably from 10% to 25% by weight.

5

ii) Silicone

The shampoo composition of the invention also comprises an insoluble, non-volatile silicone, which may be one or more 10 polyalkyl siloxanes, one or more polyalkylaryl siloxanes, or mixtures thereof. The silicone is insoluble in the aqueous matrix of the composition and so is present in an emulsified form, with the silicone present as dispersed particles.

15 Suitable polyalkyl siloxanes include polydimethyl siloxanes which have the CAFTAN designation dimethicone, having a viscosity of from 5 to 100,000 centistokes at 25°C. These siloxanes are available commercially from the General Electric Company as the Viscasil series and from Dow Corning 20 as the DC 200 series. The viscosity can be measured by means of a glass capillary viscometer as set out further in Doe Corning Corporate Test Method CTM004 July 20 1970.

Also suitable is polydiethyl siloxane.

25

The polyalkylaryl siloxanes which may be used in the compositions of the invention include polymethylphenyl polysiloxanes having a viscosity of from 15 to 65 centistokes at 25°C. The siloxanes are available 30 commercially from the General Electric Company as SF1075 methyl phenyl fluid or from Dow Corning as 556 Cosmetic Grade Fluid.

Also suitable are silicone gums, such as those described in US Pat. No. 4,152,416 (Spitzer), and on General Electric Silicone Rubber product Data Sheet SE 30, SE 33, SE 54 and SE 76. "Silicone gum" denotes polydiorganosiloxanes having 5 a molecular weight of from 200,000 to 1,000,000 and specific examples include polydimethyl siloxane polymers, polydimethyl siloxane/diphenyl/methylvinylsiloxane copolymers, polydimethylsiloxane/methylvinylsiloxane copolymers and mixtures thereof.

10

Aminofunctional silicones which have the CTFA designation amodimethicone, are also suitable for use in the compositions of the invention, as are polydimethyl siloxanes having hydroxyl end groups (which have the CTFA designation 15 dimethiconol).

The silicone materials described above are incorporated in the shampoo composition of the invention as a pre-formed aqueous emulsion. The average particle size of the silicone 20 material in this emulsion and in the shampoo composition is from 2 to 30 microns, preferably from 2 to 20 microns, more preferably 3 to 10 microns. Particle size may be measured by means of a laser light scattering technique, using a 2600D Particle Sizer from Malvern Instruments.

25

The pre-formed emulsion may be prepared by high shear mechanical mixing of the silicone and water, or by emulsifying the insoluble, non-volatile silicone with water and an emulsifier - mixing the silicone into a heated 30 solution of the emulsifier for instance, or by a combination of mechanical and chemical emulsification. A further suitable technique for preparation of the emulsions is emulsion polymerisation. Emulsion polymerised silicones as

- 9 -

such are described in US 2 891 820 (Hyde), US 3 294 725 (Findlay) and US 3 360 491 (Axon).

- Any surfactant materials either alone or in admixture may be
5 used as emulsifiers in the preparation of the pre-formed
silicone emulsions. Suitable emulsifiers include anionic,
cationic and nonionic emulsifiers. Examples of anionic
emulsifiers are alkylarylsulphonates, e.g., sodium
dodecylbenzene sulphonate, alkyl sulphates e.g., sodium,
10 lauryl sulphate, alkyl ether sulphates, e.g., sodium lauryl
ether sulphate nEO, where n is from 1 to 20 alkylphenol
ether sulphates, e.g., octylphenol ether sulphate nEO where
n is from 1 to 20, and sulphosuccinates, e.g., sodium
dioctylsulphosuccinate.
- 15 Examples of nonionic emulsifiers are alkylphenol
ethoxylates, e.g., nonylphenol ethoxylate nEO, where n is
from 1 to 50, alcohol ethoxylates, e.g., lauryl alcohol nEO,
where n is from 1 to 50, ester ethoxylates, e.g.,
20 polyoxyethylene monostearate where the number of oxyethylene
units is from 1 to 30.

- Typically, a pre-formed emulsion will contain around 50% of
silicone. Pre-formed emulsions are available from suppliers
25 of silicone oils such as Dow Corning, General Electric,
Union Carbide, Wacker Chemie, Shin Etsu, Toshiba, Toyo
Beauty Co, and Toray Silicone Co. Examples are the material
sold as DC-1310 by Dow Corning, and the materials sold as X-
52-1086, X-52-2127 and X-52-2112 by Shin-Etsu.

- 30 The compositions of the invention typically contain from
0.01 to 20% by weight, preferably from 0.1 to 10%, more
preferably from 0.25 to 3% by weight of insoluble, non-

- 10 -

volatile silicone. If less than 0.01% by weight is present in the composition, little conditioning benefit is observed, and if more than 20% by weight is present, the hair will appear greasy.

5

The aqueous pre-formed emulsion may be incorporated into the shampoo composition in an amount of from 0.02 to 40% by weight, preferably from 0.2 to 20% by weight.

10 The exact quantity of emulsion will of course depend on the concentration of the emulsion, and should be selected to give the desired quantity of insoluble, non-volatile silicone, in the final composition.

15 iii) Cationic Conditioning Polymer

A further component of hair treatment compositions of the invention is a cationic conditioning polymer.

20 The polymer may be a homopolymer or be formed from two or more types of monomers. The molecular weight of the polymer will generally be between 5 000 and 10 000 000, typically at least 10 000 and preferably in the range 100 000 to about 2 000 000. The polymers will have cationic nitrogen
25 containing groups such as quaternary ammonium or protonated amino groups, or a mixture thereof.

The cationic charge density is suitably at least 0.1 meq/g, preferably above 0.8 or higher. The cationic charge density
30 should not exceed 3 meq/g. It is preferably less than 2 meq/g. The charge density can be measured using the Kjeldahl method and should be within the above limits at the

- 11 -

desired pH of use, which will in general be from about 3 to 9 and preferably between 4 and 8.

The cationic nitrogen-containing group will generally be
5 present as a substituent on a fraction of the total monomer units of the cationic conditioning polymer. Thus when the polymer is not a homopolymer it can contain spacer non-cationic monomer units. Such polymers are described in the CTFA Cosmetic Ingredient Directory, 3rd edition.

10

Suitable cationic conditioning polymers include, for example, copolymers of vinyl monomers having cationic amine or quaternary ammonium functionalities with water soluble spacer monomers such as (meth)acrylamide, alkyl and dialkyl
15 (meth)acrylamides, alkyl (meth)acrylate, vinyl caprolactone and vinyl pyrrolidine. The alkyl and dialkyl substituted monomers preferably have C1-C7 alkyl groups, more preferably C1-3 alkyl groups. Other suitable spacers include vinyl esters, vinyl alcohol, maleic anhydride, propylene glycol
20 and ethylene glycol.

The cationic amines can be primary, secondary or tertiary amines, depending upon the particular species and the pH of the composition. In general secondary and tertiary amines,
25 especially tertiary, are preferred.

Amine substituted vinyl monomers and amines can be polymerized in the amine form and then converted to ammonium by quaternization.

30

Suitable cationic amino and quaternary ammonium monomers include, for example, vinyl compounds substituted with dialkyl aminoalkyl acrylate, dialkylamino alkylmethacrylate,

- 12 -

monoalkylaminoalkyl acrylate, monoalkylaminoalkyl methacrylate, trialkyl methacryloxyalkyl ammonium salt, trialkyl acryloxyalkyl ammonium salt, diallyl quaternary ammonium salts, and vinyl quaternary ammonium monomers
5 having cyclic cationic nitrogen-containing rings such as pyridinium, imidazolium, and quaternized pyrrolidine, e.g., alkyl vinyl imidazolium, and quaternized pyrrolidine, e.g., alkyl vinyl imidazolium, alkyl vinyl pyridinium, alkyl vinyl pyrrolidine salts. The alkyl portions of these monomers are
10 preferably lower alkyls such as the C₁-C₃ alkyls, more preferably C₁ and C₂ alkyls.

Suitable amine-substituted vinyl monomers include dialkylaminoalkyl acrylate, dialkylaminoalkyl methacrylate,
15 dialkylaminoalkyl acrylamide, and dialkylaminoalkyl methacrylamide, wherein the alkyl groups are preferably C₁-C₃, hydrocarbyls, more preferably C₁-C₃, alkyls.

The cationic conditioning polymers can comprise mixtures of
20 monomer units derived from amine- and/or quaternary ammonium-substituted monomer and/or compatible spacer monomers.

Suitable cationic conditioning polymers include, for example: copolymers of 1-vinyl-2-pyrrolidine and 1-vinyl-3-methyl-imidazolium salt (e.g., Chloride salt) (referred to in the industry by the Cosmetic, Toiletry, and Fragrance Association, "CTFA". as Polyquaternium-16) such as those commercially available from BASF Wyandotte Corp.
25 (Parsippany, NJ, USA) under the LUVIQUAT tradename (e.g., LUVIQUAT FC 370); copolymers of 1-vinyl-2-pyrrolidine and dimethylaminoethyl methacrylate (referred to in the industry by CTFA as Polyquaternium-11) such as those commercially

- 13 -

from Gar Corporation (Wayne, NJ, USA) under the GAFQUAT trademark (e.g., GAFQUAT 755N); cationic diallyl quaternary ammonium-containing polymer including, for example, dimethyldiallylammonium chloride homopolymer and copolymers 5 of acrylamide and dimethyldiallylammonium chloride, referred to in the industry (CTFA) as Polyquaternium 6 and Polyquaternium 7, respectively; mineral acid salts of amino-alkyl esters of homo-and co-polymers of unsaturated carboxylic acids having from 3 to 5 carbon atoms, as 10 described in U.S. Patent 4,009,256; and cationic polyacrylamides as described in our copending UK Application No. 9403156.4 (WO95/22311).

Other cationic conditioning polymers that can be used 15 include polysaccharide polymers, such as cationic cellulose derivatives and cationic starch derivatives.

Cationic polysaccharide polymer materials suitable for use in compositions of the invention include those of the 20 formula:

R¹

25 A-O(R-N⁺-R³X⁻)

R²

wherein: A is an anhydroglucose residual group, such as 30 starch or cellulose anhydroglucose residual,
R is an alkylene oxyalkylene, polyoxyalkylene, or hydroxyalkylene group, or combination thereof,

- 14 -

R¹, R² and R³ independently are alkyl, aryl, alkylaryl, arylalkyl, alkoxyalkyl, or alkoxyaryl groups, each group containing up to about 18 carbon atoms, and the total number of carbon atoms for each cationic moiety (i.e., the sum of 5 carbon atoms in R¹, R² and R³) preferably being about 20 or less, and X is an anionic counterion, as previously described.

Cationic cellulose is available from Amerchol Corp. (Edison, 10 NJ, USA) in their Polymer JR (trade mark) and LR (trade mark) series of polymers, as salts of hydroxyethyl cellulose reacted with trimethyl ammonium substituted epoxide, referred to in the industry (CTFA) as Polyquaternium 10. Another type of cationic cellulose includes the polymeric 15 quaternary ammonium salts of hydroxyethyl cellulose reacted with lauryl dimethyl ammonium-substituted epoxide, referred to in the industry (CTFA) as Polyquaternium 24. These materials are available from Amerchol Corp. (Edison, NJ, USA) under the tradename Polymer LM-200.

20

Other cationic conditioning polymers that can be used include cationic guar gum derivatives, such as guar hydroxypropyltrimonium chloride (Commercially available from Celanese Corp. in their Jaguar trademark series). Other 25 materials include quaternary nitrogen-containing cellulose ethers (e.g., as described in U.S. Patent 3,962,418), and copolymers of etherified cellulose and starch (e.g., as described in U.S. Patent 3,958,581).

30 Preferably the cationic conditioning polymer is selected from the group comprising cationic polyacrylamides, hydroxyalkyl cellulose ethers and cationic guar derivatives. Particularly preferred are Jaguar C13S with a cationic

- 15 -

charge density of 0.8meq/g. Jaguar C13S is guar hydroxypropyltrimonium chloride. Other particularly suitable materials include Jaguar C15, Jaguar C17 and Jaguar C16 and Jaguar C162. A preferred cellulose ether is Polymer 5 JR400.

As discussed above, we have surprisingly found that in compositions of our invention, inclusion of cationic polymer enhances the wet properties of the shampoo without 10 subsequent dry hair sensory negatives such as slippy feel leading to styling difficulties.

Advantageously, the amount of cationic conditioning polymer present in compositions on the invention is not governed by 15 the requirement that these materials should act as a deposition aid for the silicone component. The particular level appropriate in compositions of the present invention is dependent on the particular surfactant system employed. Generally, the level can vary from 0.01 to 3% by weight.

20 The invention is also directed to the use, in a shampoo composition comprising a major proportion of a surfactant, of an aqueous pre-formed emulsion of an insoluble, non-volatile silicone having an average particle size in the 25 emulsion and in the shampoo composition of from 2 to 30 microns for imparting improved conditioning benefit to hair from the shampoo composition.

The shampoo composition of the invention may further 30 comprise from 0.1 to 5 % of a silicone suspending agent. Examples are polyacrylic acids, cross linked polymers of acrylic acid, copolymers of acrylic acid with a hydrophobic monomer, copolymers of carboxylic acid- containing monomers

and acrylic esters, cross-linked copolymers of acrylic acid and acrylate esters, heteropolysaccharide gums and crystalline long chain acyl derivatives. The long chain acyl derivative is desirably selected from ethylene glycol 5 stearate, alkanolamides of fatty acids having from 16 to 22 carbon atoms and mixtures thereof. Ethylene glycol distearate and Polyethylene glycol 3 distearate are preferred long chain acyl derivatives. Polyacrylic acid is available commercially as Carbopol 420, Carbopol 488 or 10 Carbopol 493. Polymers of acrylic acid cross-linked with a polyfunctional agent may also be used, they are available commercially as Carbopol 910, Carbopol 934, Carbopol 940, Carbopol 941 and Carbopol 980. An example of a suitable 15 copolymer of a carboxylic acid containing a monomer and acrylic acid esters is Carbopol 1342. All Carbopol materials are available from Goodrich and Carbopol is a trade mark.

Suitable cross linked polymers of acrylic acid and acrylate 20 esters are Pemulen TR1 or Pemulen TR2. A suitable heteropolysaccharide gum is xanthan gum, for example that available as Kelzan mu.

Depending on the type of composition employed, one or more 25 additional ingredients conventionally incorporated into hair treatment compositions may be included in compositions of the invention. Such additional ingredients include styling agents, such as resins and hair-setting polymers, perfumes, dyes, buffering or pH adjusting agents, viscosity modifiers, 30 opacifiers, pearlescers, preservatives, antibacterial agents, antidandruff agents, foam boosters, proteins, moisturising agents, herb or other plant extracts and other natural ingredients.

- 17 -

The invention is further illustrated by way of the following non-limitative examples:

EXAMPLES

- 5 Shampoo compositions were prepared by mixing the following components in the amounts stated:

Example 1

		<u>% wt</u>
10	-----	
	Ammonium lauryl sulphate	14.0
	Cocamidopropyl betaine	2.0
15	Silicone Oil ⁽¹⁾	2.0
	Carbopol 980 ⁽²⁾	0.4
	Jaguar C13S ⁽³⁾	0.2
	Preservative, perfume, viscosity modifier	q.s.
20	Water	to 100.0

25 ⁽¹⁾ Silicone oil was included as DC-1310 from Dow Corning Ltd., an emulsion of dimethicone (60,000 cs) in nonionic surfactant.

30 ⁽²⁾ Carbopol 980 is a cross-linked polyacrylate available from B F Goodrich.

⁽³⁾ Jaguar C13S is guar hydroxypropyltrimonium chloride available from Celanese Corp.

- 19 -

Example 2

		<u>% wt</u>
	Ammonium lauryl sulphate	14.0
5	Cocamidopropyl betaine	2.0
	Silicone oil ⁽¹⁾	2.0
	Carbopol 980	0.4
	Ethylene glycol monostearate	1.5
	Jaguar C13S	0.2
10	Preservative, perfume, viscosity modifier	q.s.
	Water	to 100.0

⁽¹⁾ as Example 1

15 Example 3

		<u>% wt</u>
	Sodium lauryl ether sulphate 2EO	12.0
	Cocamidopropyl betaine	4.0
20	Silicone oil ⁽¹⁾	0.5
	Carbopol 980	0.4
	Jaguar C13S	0.03
	Ethylene glycol monostearate	1.5
	Preservative, perfume, viscosity modifier	q.s.
25	Water	to 100.0

⁽¹⁾ as Example 1

- 30 All the shampoos of Examples 1-3 are prepared using a simple cold process whereby all the ingredients are mixed using a paddle stirrer.

- 20 -

The silicone particles in the emulsion have a mean particle size of 7.7 microns and remain the same in the shampoo composition.

5 Example 4 and Comparative Example A

Evaluation of the wet/dry conditioning performance of shampoo formulations incorporating pre-emulsified silicone of different particle sizes and with or without cationic polymer

10

Test Shampoos

Shampoo compositions were prepared containing ingredients as shown in the following Table. Example 4 is a formulation
15 according to the present invention. Comparative Example A is a formulation in which the pre-emulsified large particle size silicone of Example 4 has been substituted by an equivalent amount of a pre-emulsified smaller particle size silicone as described in the prior art. Control 1 is the
20 formulation of Example 4 minus cationic polymer (Jaguar C-13-S). Control 2 is the formulation of Comparative Example A minus cationic polymer (Jaguar C-13-S).

- 21 -

Ingredient	Example 4	Control 1	Comp.Ex.A	Control 2
SLES (100%)	14	14	14	14
CAPB (100%)	2	2	2	2
Jaguar C-13-S	0.2	-	0.2	-
DC-1310 (60%)	3.3	3.3	-	-
BY22-026 (50%) ⁽¹⁾	-	-	4	4
Carbopol 980	0.4	0.4	0.4	0.4
Formalin	0.1	0.1	0.1	0.1
water	to 100	to 100	to 100	to 100

(1) BY22-026 is an emulsion of 60,000cs dimethicone with a
 5 particle size of 0.5microns, ex Toray Silicone Co.

Conditioning performance

10 The conditioning performance of each of the above test shampoos was evaluated as follows:

15 7g of hair in the form of a switch was worked in 0.7g of the test product, lathered for 30 seconds, and rinsed with water. The procedure was repeated once. Three switches of

- 22 -

hair were prepared for each product to be evaluated. The evaluation of conditioning performance was carried out by twelve trained panellists as a paired comparison test and significant differences at greater than 95% confidence were
5 assessed.

Results

- Results are shown in the following table, with each row
10 including entries for two shampoos being compared in the form of relative allocation of a total score of 100, a higher score indicating preference for that member of the pair.

15 The two attributes assessed by the panellists were (a) ease of wet combing and (b) ease of dry combing.

The results were as follows:

	Ease of wet comb	Ease of dry comb
Example 4 versus Control 1	70	26
Comp.Ex.A versus Control 2	72	80

20

The results show that:

Example 4 gives significantly greater ease of wet combing than Control 1 on ease of wet combing

25

Control 1 gives significantly greater ease of dry combing than Example 4

- 23 -

Comparative Example A gives significantly greater ease of wet and dry combing than Control 2

5 Conclusions

Example 4 according to the invention shows that inclusion of cationic polymer increases the ease of wet combing imparted by the formulation and reduces the ease of dry combing.

10

Comparative Example A shows that if the large particle size pre-emulsified silicone of the invention is substituted with a smaller particle size pre-emulsified silicone according to the prior art, then a selective increase in wet combing

15 performance is not observed. In contrast to the Example of the invention, Comparative Example A shows that inclusion of cationic polymer increases the ease of wet combing imparted by the formulation but increases the ease of dry combing (as expressed by panellist preference) to a greater extent.

20

CLAIMS

1. An aqueous shampoo composition comprising, in addition to water:

5

i) at least one surfactant chosen from anionic, nonionic, zwitterionic or amphoteric surfactants or mixtures thereof;

10 ii) emulsified particles of an insoluble, nonvolatile

silicone;

15 iii) a soluble cationic hair conditioning polymer having a cationic charge density of about +3.0 meq/gram or less,

in which the emulsified particles of insoluble, nonvolatile silicone are incorporated into the shampoo composition as a preformed aqueous emulsion having an average silicone particle size in the emulsion and in the shampoo composition of from 2 to 30 microns.

20

2. A shampoo composition according to claim 1, in which the at least one surfactant is present in an amount of from 0.1 to 50% by weight of the composition.

25

3. A shampoo composition according to claim 1, in which the anionic surfactant is selected from the group consisting of sodium lauryl sulphate, triethanolamine lauryl sulphate, triethanolamine monolauryl phosphate, sodium lauryl ether sulphate 2EO, sodium lauryl ether sulphate 3EO, ammonium lauryl sulphate, ammonium lauryl ether sulphate 1EO, ammonium lauryl ether sulphate 2EO, ammonium lauryl ether sulphate 3EO and mixtures thereof.

4. A shampoo composition according to claim 1, in which the amphoteric surfactant is selected from the group consisting of cocamidopropylbetaine, lauryl betaine and sodium cocamphopropionate.

5. A shampoo composition according to claim 1, in which the insoluble, non volatile silicone is present in an amount 10 of from 0.01 to 20% by weight of the composition.

6. A shampoo composition according to claim 1, in which the insoluble, non volatile silicone is selected from the group consisting of polydimethylsiloxanes and 15 polydimethylsiloxanes having hydroxyl end groups.

7. A shampoo composition according to claim 1, in which the cationic hair conditioning polymer is present in an amount of 0.01 to 3% by weight of the composition.

20 8. A shampoo composition according to claim 1, in which the cationic hair conditioning polymer is selected from the group consisting of cationic derivatives of guar gum, cationic cellulose ether derivatives and cationic 25 polyacrylamides.

9. A method of making the aqueous shampoo composition of Claim 1, comprising mixing together water, the surfactant, the cationic conditioning polymer and a preformed aqueous 30 emulsion of the silicone, wherein the silicone in the emulsion and in the shampoo composition has an average particle size of from 2 to 30 microns.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 97/03718

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 6 A61K7/06 A61K7/50

According to International Patent Classification(IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
 IPC 6 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, X	<p>WO 96 32919 A (PROCTER & GAMBLE) 24 October 1996 see page 3, line 9 - page 4, line 6 see page 14, line 5 - page 15, line 34 see page 16, line 25 - page 17, line 9 see page 17, line 25 - page 18, line 9 see page 28, line 23 - page 33, line 26 see claims 1-10; examples ---</p>	1-9
X	<p>WO 92 10162 A (PROCTER & GAMBLE) 25 June 1992 cited in the application see page 5, line 25 - page 6, line 14 see page 26, line 25 - page 27, line 5 see page 31, line 25-35 see claims 1-9; examples --- -/-</p>	1-9

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

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1

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

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X	WO 92 10163 A (PROCTER & GAMBLE) 25 June 1992 see page 28, line 11-26 see page 33, line 6-18; examples 1-5 ----	1-9
X	WO 95 09599 A (UNILEVER PLC ;UNILEVER NV (NL)) 13 April 1995 see page 2, line 17 – page 3, line 10 see page 11, line 13-30 see page 12, line 1 – page 13, line 23 see claims 1,2,4-10; examples 1-3 ----	1-9
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A	WO 94 03152 A (UNILEVER PLC ;UNILEVER NV (NL)) 17 February 1994 see page 15, line 21 – page 16, line 2; claims 1-9; examples 1,2 -----	1-9

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